- 1. A composition of matter comprising
 - a. integrin/adhesion antagonist peptide; and
 - b. a vehicle.
- 5 2. A composition of the formula

$$(X^1)_a - F^1 - (X^2)_b$$

and multimers thereof/wherein:

F¹ is an Fc domain;

 X^1 and X^2 are each independently selected from -(L¹)_c-P¹, -

$$(L^1)_c - P^1 - (L^2)_d - P^2$$
, $-(L^1)_c - P^1 - (L^2)_d - P^2 - (L^3)_e - P^3$, and $-(L^1)_c - P^1 - (L^2)_d - P^2 - (L^3)_e - P^3 - (L^3)_e - (L^3$

 $(L^4)_f - P^4$

P¹, P², P³, and P⁴ are each independently sequences of integrin/adhesion antagonist peptides;

 L^1 , L^2 , L^3 and L^4 are each independently linkers; and

a, b, c, d, e, and f are each independently 0 or 1, provided that at least one of a and b is 1.

The composition of matter of Claim 1 of the formulae

X¹-

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4. The composition of matter of Claim 3 of the formula

5. The composition of matter of Claim 3 of the formula

$$F^{1}-(L^{1})_{c}-P^{1}-(L^{2})_{d}-P^{2}$$

- 25 6. The composition of matter of Claim 2, wherein F¹ is an Fc domain.
 - 7. The composition of matter of Claim wherein F¹ is an IgG Fc domain.
 - 8. The composition of matter of Claim 2 wherein F¹ is an IgG1 Fc domain.

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- 9. The composition of matter of Claim 2 wherein F¹ comprises the sequence of SEQ ID NO: 2.
- 10. The composition of matter of Claim 2 wherein X^1 and X^2 comprise one or more sequences selected from SEQ ID NOS: 7 to 21.
- The composition of matter of Claim 2 wherein the composition of matter comprises one or more sequences selected from SEQ ID NOS: 22 to 94.
 - 12. The composition of matter of Claim 2, wherein the composition of matter comprises one or more sequences selected from SEQ ID NOS: 7 and 9 to 16.
 - The composition of matter of Claim 2 wherein the composition of matter comprises one or more sequences selected from Tables 3, 4, 5, and 6 (SEQ ID NOS: 22 to 94, 128 to 137).
 - 14. ADNA encoding a composition of matter of any of Claims 6 to 13.
- 15 15. An expression vector comprising the DNA of Claim 14.
 - 16. A host cell comprising the expression vector of Claim 15.
 - 17. The cell of Claim 16, wherein the cell is an <u>E. coli</u> cell.
 - 18. A process for preparing a pharmacologically active compound, which comprises
- a) selecting at least one randomized integrin/adhesion antagonist peptide; and
 - b) preparing a pharmacologic agent comprising at least one Fc domain covalently linked to at least one amino acid sequence of the selected peptide or peptides.
- 25 19. The process of Claim 18, wherein the peptide is selected in a process comprising one or more techniques selected from yeast-based screening, rational design, protein structural analysis, screening of a phage display library, an <u>E. coli</u> display library, a ribosomal library, or a chemical peptide library.

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- 20. The process of Claim 18, wherein the preparation of the pharmacologic agent is carried out by:
 - a) preparing a gene construct comprising a nucleic acid sequence encoding the selected peptide and a nucleic acid sequence encoding an Fc domain; and
 - b) expressing the gene construct.
- 21. The process of Clarm 18, wherein the gene construct is expressed in an <u>E. coli</u> cell.
- 22. The process of Claim 18 wherein the Fc domain is an IgG Fc domain.
- 23. The process of Claim 18, wherein the vehicle is an IgG1 Fc domain.
- 24. The process of Claim 18, wherein the vehicle comprises the sequence of SEQ ID NO: 2.
- A composition of matter comprising an amino acid sequence selected from SEQ ID NOS: 132 to 137.